

POLYMERIC COMPOSITIONS

[0001] The present invention relates to improved formulations using certain polymer precursors which have advantageous properties for example for use as inks e.g. for flexographic applications.

BACKGROUND OF THE INVENTION

[0002] A new class of low molecular weight and/or functional acrylic polymers have recently been discovered and are described in, for example, U.S. Serial No. 60/215,721 (abandoned), co-pending U.S. patent application Serial No. 09/886,192 (Rohm & Haas), and U.S. patents US 4,414,370 (S C Johnston), U.S. 4,529,787 (S C Johnston), U.S. 4,546,160 (S C Johnston), U.S. 5,010,166 (S C Johnston), U.S. 5,475,073 (Arco), U.S. 5,525,693 (Arco), U.S. 5,646,213 (Arco) and U.S. 5,646,225 (Arco), the subject matter of which is incorporated by reference herein

[0003] The applicant has surprisingly discovered that when such oligomers are used to prepare formulations for example for radiation curing in the absence of undesirable species such as volatile organic compounds (VOCs), such formulations (e.g. radiation curable inks) exhibit advantageous properties.

BRIEF SUMMARY OF THE INVENTION

[0004] Therefore broadly in accordance with the present invention there is provided a radiation-curable formulation substantially-free of volatile organic compounds (VOCs), the formulation comprising:

- (a) from about 5% to about 70% by weight of an a monomer diluent comprising at least one activated unsaturated moiety;
- (b) optionally up to about 10% by weight a photo-initiator; and

(c) from about 30% to about 95% by weight of at least one polymer and/or oligomer obtained and/or obtainable from a monomer comprising at least one activated unsaturated moiety,

where the formulation and/or components thereof exhibit at least one of the following properties:

(w) the polymer component (c) has a solubility parameter (heat of fusion per unit volume) of more than about $15 \text{ J}^{1/2} \text{cm}^{-3/2}$;

(x) the polymeric component (c) has a number average molecular weight (M_n) of less than about 15,000 daltons;

(y) the formulation has a viscosity of up to about 10,000 centipoise and no more than 50% of diluent; and/or

(z) the polymeric component (c) having a dispersing power (Θ) as defined herein of up to about 1.0 .

DETAILED DESCRIPTION OF THE INVENTION

[0005] The monomer used to form polymeric component (c) may be the same as or different from the diluent monomer(s) used in component (a) .

[0006] A solubility parameter (SP) is a means of quantifying the way that "like dissolves like". SP can be derived from a cohesive energy density defined as:

$$\text{Cohesive Energy Density (CED)} = \frac{\Delta E}{V} = \frac{\Delta H - RT}{V}, \text{ where}$$

ΔE = Energy required to vaporize 1 cc of solvent;

V = Molar volume (i.e. molecular weight /density);

ΔH = Latent heat of vaporization;

R = Gas constant

T = Temperature

The SP can thus be based on three components (following the known approach used by van Krevelen) namely a dispersive (non-polar) component, a polar component and a hydrogen bonding component. Thus the SP can be defined as

$$\text{Solubility Parameter} = \sqrt{\text{CED}} = \sqrt{\text{CED}_{\text{Dispersive}}^2 + \text{CED}_{\text{Polar}}^2 + \text{CED}_{\text{H-Bonding}}^2}$$

[0007] The SP can be used to measure resin compatibility, resin solubility and/or resin and diluent/solvent compatibility. SP can be determined indirectly by testing solubility in solvents with known solubility parameters, then plotting on SP chart (e.g., H-bonding vs. Polar). To do this the resin is dissolved in solvent, titrated to a cloud point with non-solvent, and then the solubility parameter can be calculated on solvent blend composition (e.g., KB value). One can also calculate SP from the molecular structure of a resin, for example from structural groups such as CH₃-, phenyl, C=O, -OH, CH₂=, etc.

[0008] Preferably for oligomers used to formulate preferred inks of the invention as described and exemplified herein, the oligomer exhibits a solubility parameter of from about 15 J^{1/2}cm^{-3/2} to about 30 J^{1/2}cm^{-3/2}; more preferably from about 18 J^{1/2}cm^{-3/2} to about 25 J^{1/2}cm^{-3/2}.

[0009] Preferably the number average molecular weight (M_n) of the polymer component (c) is from about 200 to about 20,000 daltons, more preferably from about 500 to about 15,000 daltons, most preferably from about 1000 to about 10,000 daltons.

[0010] Preferred formulations of the invention have a viscosity of up to about 2,000 centipoise per g of polymeric component (c) at a temperature of 25°C.

[0011] The dispersing power (denoted herein by Θ) of a functional polymer is given as a product of two parameters ε and ρ divided by the average molecular weight of the polymer to give value of the dispersing strength of the polymer per unit weight, i.e.

$$\Theta = \epsilon \times \rho / M_w$$

Preferably Θ is from about 1×10^{-5} to about 0.8, more preferably from about 1×10^{-3} to about 0.5; most preferably from about 0.01 to about 0.1 DP (equivalents) / (mol x daltons).

[0012] ϵ is a measure of the ability of a polymer to disperse a polar substance (such as a colorant) in a suitable fluid and is the product of the average electric dipole moment μ_a and to the average polarizability α_a of all the polar substituent groups on the polymer, thus

$$\epsilon = \mu_a \times \alpha_a$$

ϵ is measured in units of DP ($1 \text{ DP} = 3.336 \times 10^{-30} \text{ C}\cdot\text{m} \times 1 \times 10^{-24} \text{ cm}^3 = 3.336 \times 10^{-58} \text{ C}\cdot\text{m}^4$) and preferably is less than about 400 DP ($\text{C}\cdot\text{m}$ = coulomb meter).

[0013] ρ is a measure of the concentration of all polar substituents on a polymer measured in units of equivalents per mol (of polymer); and is preferably less than about 100 equivalents per mol.

[0014] M_w is the weight average molecular weight of the polymer (measured by GPC / MS using a polystyrene standard) and is preferably less than about 25,000 daltons.

[0015] An electric dipole consists of two electric charges $-q$ and $+q$ separated by a distance d . It is represented by a vector directed from the negative to the positive charge. The magnitude of this vector is given by "qd" and is called the electric dipole moment, which is conveniently measured herein in Debye units (D) where $1 \text{ D} = 3.336 \times 10^{-30} \text{ C}\cdot\text{m}$. A polar group is a chemical species having a permanent dipole moment (also denoted herein by " μ ") which exists independently of any applied electric field. This occurs whenever the mean center of the negative charge on the species is different from the mean center of the positive charge on

that species. All values for the (scalar) electric dipole moment quoted herein are those obtained when measured in the gas phase by standard techniques well known in the art.

[0016] The electric dipole polarizability of a chemical species is equal to the electrical moment of a dipole induced in that species divided by the field intensity. Conveniently the electric dipole polarizability measured as an average over the chemical species' (e.g. molecule's) ground state is given herein in units of $1 \times 10^{-24} \text{ cm}^3$ (which is denoted herein by the abbreviation "P"). Polarizability can be considered a measure of the ability of a species to exhibit a whole or partial electric charge and is also denoted herein by " α ".

[0017] Thus the parameter μ_a is a measure of the overall polarity of the polymer component (c) and is a scalar quantity independent of the sign of the charge and α_a is a reflection of the readiness of the polymer component (c) to hold charge and both together give a measure of the ease of which a given amount of polymer will disperse a material containing polar groups such as a common colorants which contain polar chromophores such as azo groups and/or phthalocyanine rings.

[0018] The polymeric component (c) may comprise those polar substituents thereon which on average have exhibit an electric dipole moment (μ_a) of from about 0.01 D to about 15 D and/or an average electric polarizability (α_a) of from about 1 to about 25 P; thus preferably ϵ may be from about 0.01 to about 375 DP.

[0019] Preferably μ_a is from about 0.01 D to about 10 D, more preferably from about 0.5 D to about 7 D; most preferably from about 1 D to about 5 D.

Preferably α_a is from about 2 P to about 20 P more preferably about 4 P to about 17 P; most preferably about 5 P to about 15 P.

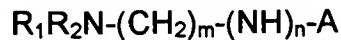
[0020] Preferably the density of polar substituents (p) is from about 1 to about 100, more preferably from about 2 to about 50, most preferably from about 3 to about 20 equivalents per mol of polymer.

[0021] Preferably the molecular weight (Mw) of the polymer component (c) is from about 1,000 to about 25,000 daltons, more preferably from about 1,500 to about 20,000 daltons, most preferably from about 2000 to about 15,000 daltons.

[0022] Preferably component (a) and/or (c) comprise an oligomer, more preferably which is polymerisable by radiation, most preferably which comprises at least one activated unsaturated moiety as defined herein.

[0023] Component (a) and/or (c) may conveniently be selected from one or more of the specific polymer types I to IX as described herein.

[0024] Polymers of type I comprise types A, B and C as described and exemplified herein, preferably comprise those described in the co-pending application USSN 60/215,721 or 09/886,192 (Rohm & Haas), the contents of which are hereby incorporated by reference; more preferably comprise semi-telechelic nitrogen-functional oligomers of formula



in which

A is the residue of a macromonomer bearing at least one pendant ethylenic unsaturation with a degree of polymerization from about 2 to about 50;

n is 0 or 1;

m is from 0 to 18

R₁ and R₂ are both independently selected from H, and optionally substituted C₁₋₁₈hydrocarbo;

Preferably when n is 0, m is 0 and when n is 1; m is from 0 to 18, more preferably from 1 to 18.

[0025] Preferably R₁ and R₂ are both independently selected from H, and C₁₋₁₂branched, unbranched and cyclic alkyl, phenyl and substituted phenyl where R₁ and R₂ are other than both H;

Preferably R₁R₂N- forms a cyclic group.

[0026] Polymers of type II comprise those described in US 4,414,370 (S.C. Johnston) the contents of which are hereby incorporated by reference, more preferably comprise low molecular weight vinylic polymers of at least one monoalkenyl aromatic monomer and at least one acrylic monomer having a dispersion index less than about 2 and a number average molecular weight from about 1000 to 6000 obtained and/or obtainable by a bulk polymerisation process which comprises the steps of continuously:

- (a) charging a mixture of vinylic monomers into a continuous stirred reactor zone containing a molten resin mixture of unreacted vinylic monomers and said vinylic polymer product;
- (b) maintaining the molten resin mixture at a reaction temperature from about 235°C. to 310°C.; and
- (c) maintaining a flow rate through said reaction zone sufficient (1) to provide a residence time of said charged vinylic monomer mixture in said reaction zone of at least about 2 minutes to provide a reaction product; and (2) to maintain a predetermined level of reaction mixture in said reaction zone.

[0027] Polymers of type III comprise those described in US 4,529,787 (S.C. Johnston) the contents of which are hereby incorporated by reference, more

preferably comprise low molecular weight vinylic polymer having a narrow molecular weight distribution and a low chromophore content, the polymer obtained and/or obtainable by a bulk polymerization process comprising the steps of continuously:

- (a) charging into a continuous mixed reactor zone containing a molten resin mixture;
- (i) a mixture of vinyl monomers comprising at least one monoalkenyl aromatic monomer and at least one acrylic monomer;
- (ii) a polymerization initiator in amounts to provide a molar ratio of said initiator to said mixture of vinyl monomers from about 0.0005:1 to 0.04:1;
- (iii) from about 0 to 25 percent based on the weight of vinyl monomers of a reaction solvent, wherein said molten resin mixture comprises unreacted vinylic monomers and the vinylic polymer product;
- (b) maintaining a flow rate through said reaction zone sufficient to:
- (i) provide a residence time of said charged vinylic monomer mixture in said reaction zone of from about two minutes to one hour; and
- (ii) maintain a predetermined level of reaction mixture in said reaction zone, and;
- (c) maintaining the molten resin mixture at a reaction temperature within the range from about 180°C. to 270°C. sufficient to provide accelerated conversion to a readily processable, uniform, concentrated polymer product having a number average molecular weight of 500 to 6000, a polydispersity of less than about 2.5 and a distribution index of less than about 4.5

[0028] Polymers of type IV comprise those described in US 4,546,160 (S.C. Johnston) the contents of which are hereby incorporated by reference; more preferably comprise high solids, non-styrenic acrylic polymers having a number average molecular weight of about 1000 to 2500 a polydispersity ratio of less than about 3; a dispersion index of up to about 5 and a low chromophore content,

the polymer obtained and/or obtainable by a continuous bulk polymerization process comprising the steps of continuously:

- (a) charging into a continuous mixed reactor zone containing a molten resin mixture consisting essentially of;
 - (i) at least one acrylic monomer;
 - (ii) a polymerization initiator in amounts to provide a molar ratio of said initiator to said acrylic monomer from about 0.0005:1 to 0.06:1,
 - (iii) from about 0 to 25 percent based on the weight of acrylic monomers of a reaction solvent, wherein said molten resin mixture comprises unreacted acrylic monomers and the acrylic polymer product;
- (b) maintaining a flow rate through said reaction zone sufficient to:
 - (i) provide a residence time of said charged acrylic monomer in said reaction zone of from about 1-30 minutes; and
 - (ii) maintain a predetermined level of reaction mixture in said reaction zone, and;
- (c) maintaining the molten resin mixture at an elevated temperature with the range of from about 180°C. to 270°C. sufficient to provide accelerated conversion to a readily processable, uniform, concentrated polymer product.

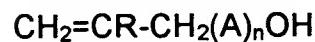
[0029] Polymers of type V comprise those described in US 5,010,166 (S.C. Johnston) the contents of which are hereby incorporated by reference; more preferably comprise graft polymers having at least one main chain and a plurality of side chains attached thereto obtained and/or obtainable by a continuous polymerization process comprising the steps of:

combining in a reaction zone an (1) addition-polymerizable monomer that is also a free-radical initiator together with an (2) ethylenically-unsaturated monomer having a nucleophilic or an electrophilic moiety at a reaction temperature that is effective for initiating addition co-polymerization of the addition-polymerizable monomer with the ethylenically-unsaturated monomer, wherein propagation by the addition co-polymerization reaction forms the main chain of the graft polymer;

while also combining in the reaction zone a polymerizable, carbonyl carbon-containing ringed molecule at a super atmospheric pressure effective, at the elevated temperature, to cause the ring portion of the ringed molecule to open in the vicinity of the carbonyl carbon thereof in response to the presence of the nucleophilic or the electrophilic moiety of the ethylenically-unsaturated monomer, for initiating ionic co-polymerization of the polymerizable ringed molecule with the nucleophilic-containing or electrophilic-containing moiety of the ethylenically-unsaturated monomer, wherein propagation by the ionic co-polymerization reaction forms the side chains of the graft polymer; and terminating both of the addition co-polymerization and the ionic co-polymerization reactions when the thus-produced graft polymer attains a predetermined number-average and/or weight-average molecule weight, the polymerization process being characterized in that each of the main-chain and side-chain propagations occurs substantially simultaneously, relative to the other, and wherein the polymerization process is further characterized in that the ionic co-polymerization reaction occurs expressly excluding utilization of a catalyst for the carbonyl carbon-containing ringed molecule.

[0030] Polymers of type V comprise low molecular weight hydroxy-functional acrylate resins optionally water reducible and/or water dispersible, the resin comprising recurring units of:

(a) an allylic alcohol (optionally in amount of about 5 to about 50% by weight), the alcohol having the general structure:



in which

n is less than about 3.0;

A is a (C_{1-12} hydrocarbylene)oxy group; and

R is selected from the group consisting of: hydrogen and C_{1-12} hydrocarbyl;

- (b) a C₁₋₂₀hydrocarbyl; vinyl aromatic; aryl acrylate and/or methacrylate monomer; (optionally in amount of about 50% to about 90% by weight);
(c) optionally, one or more ethylenically unsaturated monomers (optionally in an amount from about 1 to about 50% by weight), the monomers selected from the group consisting of vinyl aromatic monomers, unsaturated nitriles, vinyl esters, vinyl ethers, vinyl halides, vinylidene halides, unsaturated anhydrides, unsaturated dicarboxylic acids, (meth)acrylic acids; (meth)acrylates, (meth)acrylamides and conjugated dienes;

where the acrylate resin has:

- a hydroxyl number from about 15 to about 500 mg KOH/g;
an acid number within the range of about 5 to about 330 mg KOH/g;
a number average molecular weight M_n from about 500 to about 10,000 daltons;
and/or
a T_g from about -20°C. to about 50°C as measured by differential scanning calorimetry (DSC).

[0031] Polymers of type VI comprise those described in US 5,475,073 (Arco) the contents of which are hereby incorporated by reference; more preferably comprise low-molecular-weight, hydroxy-functional acrylate resins which comprise recurring units of:

- (a) a propoxylated allylic alcohol of the formula



in which

A is an oxypropylene group,

R is selected from the group consisting of hydrogen and C₁₋₅alkyl, and n, which is the average number of oxypropylene groups in the propoxylated allylic alcohol, has a value less than or equal to 2;

- (b) C₁₋₂₀alkyl or aryl acrylate or methacrylate monomer, and

(c) optionally, one or more ethylenic monomers selected from the group consisting of vinyl aromatic monomers, unsaturated nitriles, vinyl esters, vinyl ethers, vinyl halides, vinylidene halides, unsaturated anhydrides, unsaturated dicarboxylic acids, acrylic and methacrylic acids, acrylamide and methacrylamide, and conjugated dienes;

where the acrylate resin has a hydroxyl number within the range of about 50 to about 450 mg KOH/g, and a number average molecular weight within the range of about 500 to about 10,000.

[0032] Polymers of type VII comprise those described in US 5,525,693 (Arco) the contents of which are hereby incorporated by reference; more preferably comprise low molecular weight, hydroxy functional acrylate resins which comprise recurring units of:

(a) an allylic alcohol having the general structure



in which R is selected from the group consisting of hydrogen and C₁₋₅alkyl, and

(b) a C₁₋₂₀alkyl or aryl acrylate or methacrylate monomer; and

(c) optionally, one or more ethylenic monomers selected from the group consisting of vinyl aromatic monomers, unsaturated nitriles, vinyl esters, vinyl ethers, vinyl halides, vinylidene halides, unsaturated anhydrides, unsaturated dicarboxylic acids, acrylic and methacrylic acids, acrylamide and methacrylamide, and conjugated dienes;

wherein the acrylate resin has a hydroxyl number within the range of about 20 to about 500 mg KOH/g, and a number average molecular weight within the range of about 500 to about 10,000.

[0033] Polymers of type VIII comprise those described in US 5,646,213 (Arco) the contents of which are hereby incorporated by reference; more preferably comprise acrylic resins which comprise recurring units of:

(1) an allylic alcohol or a propoxylated allyl alcohol of the formula



in which

A is an oxypropylene group,

R is selected from the group consisting of hydrogen and C₁₋₅alkyl, and

n, which is the average number of oxypropylene groups in the propoxylated allylic alcohol has a value within the range of about 1 to about 2;

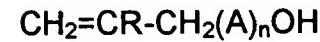
(2) a vinyl aromatic monomer; and

(3) one or more C₁₋₂₀alkyl or aryl acrylate or methacrylate monomers;

where the resin has a glass-transition temperature within the range of about -20°C. to about 50°C., a hydroxyl number within the range of about 60 to about 160 mg KOH/g, and a number average molecular weight within the range of about 1000 to about 5000

[0034] Polymers of type IX comprise those described in US 5,646,225 – (Arco) the contents of which are hereby incorporated by reference; more preferably comprise water reducible and/or water dispersible resins which comprise recurring units of:

(1) from about 5 to about 50% by weight of an allylic alcohol or a propoxylated allyl alcohol of the formula



in which

A is an oxypropylene group,

R is selected from the group consisting of hydrogen and C₁₋₅alkyl, and

n, which is the average number of oxypropylene groups in the propoxylated allylic alcohol has a value within the range of about 1 to about 2;

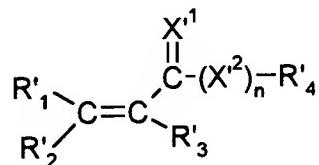
(2) from about 50 to about 90% by weight of a vinyl aromatic monomer; and

(3) from about 1 to about 50% by weight of an acrylic acid monomer;

where the resin has a number average molecular weight within the range of about 500 to about 10,000, a hydroxyl number within the range of about 15 to about 250 mg KOH/g, and an acid number within the range of about 5 to about 330 mg KOH/g.

[0035] Throughout this specification, the term "activated unsaturated moiety" "is used to denote species comprising at least one unsaturated carbon to carbon double bond in chemical proximity to at least one activating moiety. Preferably the activating moiety comprises any group which activates an ethylenically unsaturated double bond for addition thereon by a suitable electrophilic group. Conveniently the activating moiety comprises oxy, thio, (optionally organo substituted)amino, thiocarbonyl and/or carbonyl groups (the latter two groups optionally substituted by thio, oxy or (optionally organo substituted) amino). More convenient activating moieties are (thio)ether, (thio)ester and/or (thio)amide moiety(ies). Most convenient "activated unsaturated moieties" comprise an "unsaturated ester moiety" which denotes an organo species comprising one or more "hydrocarbylidene(thio)carbonyl(thio)oxy" and/or one or more "hydrocarbylidene(thio)- carbonyl(organo)amino" groups and/or analogous and/or derived moieties for example moieties comprising (meth)acrylate functionalities and/or derivatives thereof. "Unsaturated ester moieties" may optionally comprise optionally substituted generic α,β -unsaturated acids, esters and/or other derivatives thereof including thio derivatives and analogs thereof.

[0036] Preferred activated unsaturated moieties are those represented by Formula 1'.



Formula 1'

where

n' is 0 or 1,

X¹ is oxy or, thio

X² is oxy, thio or NR'₅ (where R'₅ represents H or optionally substituted organo), R'₁, R'₂, R'₃ and R'₄ each independently represent H, optionally substituents and/or optionally substituted organo groups; and all suitable isomers thereof, combinations thereof on the same species and/or mixtures thereof.

[0037] It will be appreciated that the terms "activated unsaturated moiety"; "unsaturated ester moiety" and/or Formula 1' herein may represent a discrete chemical species (such as a compound, ion, free radical, oligomer and/or polymer) and/or any part(s) thereof. Thus Formula 1' may also represent multivalent (preferably divalent) radicals. Thus the options given herein for n', X¹, X², R'₁, R'₂, R'₃, R'₄ and R'₅ also encompass corresponding bi or multivalent radicals as appropriate.

[0038] More preferred moieties of Formula 1' (including isomers and mixtures thereof) are those where n' is 1; X¹ is O; X² is O, S or NR'₅; R'₁, R'₂, R'₃, and R'₄ are independently selected from: H, optional substituents and optionally substituted C₁₋₁₀hydrocarbo, and where present R'₅ is selected from H and optionally substituted C₁₋₁₀hydrocarbo.

[0039] Most preferably n' is 1, X¹ is O; X² is O or S and R'₁, R'₂, R'₃ and R'₄ are independently H, hydroxy and/or optionally substituted C₁₋₆hydrocarbyl.

[0040] For example n' is 1, X¹ and X² are both O; and R'₁, R'₂, R'₃ and R'₄ are independently H, OH, and/or C₁₋₄alkyl.

[0041] For moieties of Formula 1' where n' is 1 and X¹ and X² are both O then: when one of (R'₁ and R'₂) is H and also R'₃ is H, Formula 1' represents an acrylate moiety, which includes acrylates (when both R'₁ and R'₂ are H) and derivatives thereof (when either R'₁ or R'₂ is not H). Similarly when one of (R'₁ and R'₂) is H and also R'₃ is CH₃, Formula 1' represents an methacrylate moiety, which includes methacrylates (when both R'₁ and R'₂ are H) and derivatives thereof (when either R'₁ or R'₂ is not H). Acrylate and/or methacrylate moieties of Formula 1' are particularly preferred.

[0042] Conveniently moieties of Formula 1' are those where n' is 1; X¹ and X² are both O; R'₁ and R'₂ are independently H, methyl or OH, and R'₃ is H or CH₃.

[0043] More conveniently, moieties of Formula 1' are those where n' is 1; X¹ and X² are both O; R'₁ is OH, R'₂ is CH₃, and R'₃ is H, and/or tautomer(s) thereof (for example of an acetoacetoxy functional species).

[0044] Most convenient unsaturated ester moieties are selected from: -OCO-CH=CH₂; -OCO-C(CH₃)=CH₂; acetoacetoxy, -OCOCH=C(CH₃)(OH) and all suitable tautomer(s) thereof.

[0045] It will be appreciated that any suitable moieties represented by Formula 1' could be used in the context of this invention such as other reactive moieties.

[0046] The terms 'optional substituent' and/or 'optionally substituted' as used herein (unless followed by a list of other substituents) signifies the one or more of following groups (or substitution by these groups): carboxy, sulpho, formyl, hydroxy, amino, imino, nitrilo, mercapto, cyano, nitro, methyl, methoxy and/or combinations thereof. These optional groups include all chemically possible combinations in the same moiety of a plurality (preferably two) of the

aforementioned groups (e.g. amino and sulphonyl if directly attached to each other represent a sulphamoyl group). Preferred optional substituents comprise: carboxy, sulpho, hydroxy, amino, mercapto, cyano, methyl, halo, trihalomethyl and/or methoxy.

[0047] The synonymous terms ‘organic substituent’ and “organic group” as used herein (also abbreviated herein to “organo”) denote any univalent or multivalent moiety (optionally attached to one or more other moieties) which comprises one or more carbon atoms and optionally one or more other heteroatoms. Organic groups may comprise organohetaryl groups (also known as organoelement groups) which comprise univalent groups containing carbon, which are thus organic, but which have their free valence at an atom other than carbon (for example organothio groups). Organic groups may alternatively or additionally comprise organyl groups which comprise any organic substituent group, regardless of functional type, having one free valence at a carbon atom. Organic groups may also comprise heterocyclyl groups which comprise univalent groups formed by removing a hydrogen atom from any ring atom of a heterocyclic compound: (a cyclic compound having as ring members atoms of at least two different elements, in this case one being carbon). Preferably the non carbon atoms in an organic group may be selected from: hydrogen, halo, phosphorus, nitrogen, oxygen, silicon and/or sulphur, more preferably from hydrogen, nitrogen, oxygen, phosphorus and/or sulphur.

[0048] Most preferred organic groups comprise one or more of the following carbon containing moieties: alkyl, alkoxy, alkanoyl, carboxy, carbonyl, formyl and/or combinations thereof; optionally in combination with one or more of the following heteroatom containing moieties: oxy, thio, sulphanyl, sulphonyl, amino, imino, nitrilo and/or combinations thereof. Organic groups include all chemically possible combinations in the same moiety of a plurality (preferably two) of the

aforementioned carbon containing and/or heteroatom moieties (e.g. alkoxy and carbonyl if directly attached to each other represent an alkoxycarbonyl group).

[0049] The term 'hydrocarbo group' as used herein is a sub-set of a organic group and denotes any univalent or multivalent moiety (optionally attached to one or more other moieties) which consists of one or more hydrogen atoms and one or more carbon atoms and may comprise one or more saturated, unsaturated and/or aromatic moieties. Hydrocarbo groups may comprise one or more of the following groups. Hydrocarbyl groups comprise univalent groups formed by removing a hydrogen atom from a hydrocarbon (for example alkyl). Hydrocarbylene groups comprise divalent groups formed by removing two hydrogen atoms from a hydrocarbon, the free valencies of which are not engaged in a double bond (for example alkylene). Hydrocarbylidene groups comprise divalent groups (which may be represented by " $R_2C=$ ") formed by removing two hydrogen atoms from the same carbon atom of a hydrocarbon, the free valencies of which are engaged in a double bond (for example alkylidene). Hydrocarbyldyne groups comprise trivalent groups (which may be represented by " $RC\equiv$ "), formed by removing three hydrogen atoms from the same carbon atom of a hydrocarbon the free valencies of which are engaged in a triple bond (for example alkylidyne). Hydrocarbo groups may also comprise saturated carbon to carbon single bonds (e.g. in alkyl groups); unsaturated double and/or triple carbon to carbon bonds (e.g. in respectively alkenyl and alkynyl groups); aromatic groups (e.g. in aryl groups) and/or combinations thereof within the same moiety and where indicated may be substituted with other functional groups

[0050] The term 'alkyl' or its equivalent (e.g. 'alk') as used herein may be readily replaced, where appropriate and unless the context clearly indicates otherwise, by terms encompassing any other hydrocarbo group such as those described herein (e.g. comprising double bonds, triple bonds, aromatic moieties (such as

respectively alkenyl, alkynyl and/or aryl) and/or combinations thereof (e.g. aralkyl) as well as any multivalent hydrocarbo species linking two or more moieties (such as bivalent hydrocarbylene radicals e.g. alkylene).

[0051] Any radical group or moiety mentioned herein (e.g. as a substituent) may be a multivalent or a monovalent radical unless otherwise stated or the context clearly indicates otherwise (e.g. a bivalent hydrocarbylene moiety linking two other moieties). However where indicated herein such monovalent or multivalent groups may still also comprise optional substituents. A group which comprises a chain of three or more atoms signifies a group in which the chain wholly or in part may be linear, branched and/or form a ring (including spiro and/or fused rings). The total number of certain atoms is specified for certain substituents for example C₁-Norgano, signifies a organo moiety comprising from 1 to N carbon atoms. In any of the formulae herein if one or more substituents are not indicated as attached to any particular atom in a moiety (e.g. on a particular position along a chain and/or ring) the substituent may replace any H and/or may be located at any available position on the moiety which is chemically suitable and/or effective.

[0052] Preferably any of the organo groups listed herein comprise from 1 to 36 carbon atoms, more preferably from 1 to 18. It is particularly preferred that the number of carbon atoms in an organo group is from 1 to 12, especially from 1 to 10 inclusive, for example from 1 to 4 carbon atoms.

[0053] As used herein chemical terms (other than IUPAC names for specifically identified compounds) which comprise features which are given in parentheses – such as (alkyl)acrylate, (meth)acrylate and/or (co)polymer - denote that that part in parentheses is optional as the context dictates, so for example the term (meth)acrylate denotes both methacrylate and acrylate.

[0054] The substituents on the repeating unit of a polymer and/or oligomer may be selected to improve the compatibility of the materials with the polymers and/or resins in which they may be formulated and/or incorporated for the uses described herein. Thus the size and length of the substituents may be selected to optimise the physical entanglement or interlocation with the resin or they may or may not comprise other reactive entities capable of chemically reacting and/or cross-linking with such other resins as appropriate.

[0055] Certain moieties, species, groups, repeat units, compounds, oligomers, polymers, materials, mixtures, compositions and/or formulations which comprise and/or are used in some or all of the invention as described herein may exist as one or more different forms such as any of those in the following non exhaustive list: stereoisomers (such as enantiomers, e.g. E and/or Z forms, diastereoisomers and/or geometric isomers); tautomers (e.g. keto and/or enol forms), conformers, salts, zwitterions, complexes (such as chelates, clathrates, crown compounds, cyptands/cryptades, inclusion compounds, intercalation compounds, interstitial compounds, ligand complexes, organometallic complexes, non-stoichiometric complexes, π -adducts, solvates and/or hydrates); isotopically substituted forms, polymeric configurations (such as homo or copolymers, random, graft and/or block polymers, linear and/or branched polymers, e.g. star and/or side branched, cross-linked and/or networked polymers, polymers obtainable from di and/or tri-valent repeat units, dendrimers, polymers of different tacticity (e.g. isotactic, syndiotactic or atactic polymers); polymorphs (such as interstitial forms, crystalline forms and/or amorphous forms), different phases, solid solutions; and/or combinations thereof and/or mixtures thereof where possible. The present invention comprises and/or uses all such forms which are effective as defined herein.

[0056] As used herein the term “volatile organic compounds” (also denoted by the abbreviation VOC) are organic compounds that readily and fully evaporate and remain in the air as gases when exposed to the atmosphere at standard temperature and pressure. VOC also comprise that range of organic compounds known as semi-volatile organic compounds that will slowly and partially evaporate when exposed to the atmosphere at standard temperature and pressure. These components tend to occur both attached to solid surfaces and as liquid droplets as well as vapor in the atmosphere. Examples of VOCs include low molecular weight species for example lower alkanes such as butane; lower alkenes such as ethylene; lower aromatics such as toluene; terpenes such as 1,8 cineole (e.g. from eucalyptus oil); and/or lower alkynes such as ethyne.

[0057] Unless the context clearly indicates otherwise, as used herein plural forms of the terms herein are to be construed as including the singular form and vice versa.

[0058] The term “comprising” as used herein will be understood to mean that the list following is non-exhaustive and may or may not include any other additional suitable items, for example one or more further feature(s), component(s), ingredient(s) and/or substituent(s) as appropriate.

[0059] The terms ‘effective’, ‘acceptable’ ‘active’ and/or ‘suitable’ (for example with reference to any process, use, method, application, preparation, product, material, formulation, compound, monomer, oligomer, polymer precursor, and/or polymers of the present invention and/or described herein as appropriate) will be understood to refer to those features of the invention which if used in the correct manner provide the required properties to that which they are added and/or incorporated to be of utility as described herein. Such utility may be direct for example where a material has the required properties for the aforementioned

uses and/or indirect for example where a material has use as a synthetic intermediate and/or diagnostic tool in preparing other materials of direct utility. As used herein these terms also denote that a functional group is compatible with producing effective, acceptable, active and/or suitable end products.

[0060] Preferred utility of the present invention comprises one or more of use as an radiation (e.g. UV and/or EB) curable ink and/or coating composition for example in lithography and/or flexography.

[0061] Other aspects of the present invention are disclosed in the claims.

[0062] The invention will now be illustrated by the following non-limiting examples. In the Examples the following abbreviations were used.

AA	acrylic acid;	APA	aminopropyl amine
APP	1-(3-aminopropyl)-2-pyrrolidone;		
BA	butyl acrylate	BDDA	1,4-butanediol diacrylate
DA	n-decyl acrylate	DMAPA	N,N-dimethylaminopropyl amine
GMA	glycidyl acrylate	HBA	4-hydroxybutyl acrylate
HEA	2-hydroxyethyl acrylate	HQ	hydroquinone
LA	lauryl acrylate	MSA	methanesulfonic acid
OA	n-octyl acrylate	ODA	a 45 / 55 (% w/w) resp. mixture of octyl acrylate & decyl acrylate.

[0063] Firstly the preparation of various known macro-monomers and oligomers are described, herein (types A to D) which include those described in U.S. application Serial No. 60/215,721 and 09/886,182, the latter co-pending, the

contents of which are incorporated by reference. These macro-monomers and oligomers may be used to prepare formulations of the invention described later.

[0064] The composition and molecular weight of the oligomers can be determined by many conventional analytical techniques. In the examples below, the oligomers were characterized by NMR. The mole ratio of the monomers in each oligomer was determined by proton NMR. The number average molecular weight (M_n) and the weight average molecular weight (M_w) were determined by gel permeation chromatography (GPC) using appropriate molecular weight standards. The average oligomer formula was calculated from the M_n and the mole ratio of monomers in the oligomer. The polydispersity of the oligomer was calculated from the ratio of M_w to M_n . The degree of polymerization of an oligomer is the number of monomer units incorporated into the oligomer backbone and is calculated from the mole ratio of monomers in the oligomer and the M_n .

A. Preparation of Macromonomers (Oligomers A.1 to A.7)

[0065] Macromonomers of type A were prepared as described below. These monomers may be used as precursors to prepare other macromonomers as described herein (e.g. type D acrylated oligomers and/or type B nitrogen functional oligomers) and/or may be used directly to prepare formulations of the present invention.

[0066] The macromonomers were prepared by a continuous high temperature polymerization process. The polymerization reactor was a 3.05 meter (10 feet) long section of stainless steel tubing having a inner diameter of 1.6 mm (1/16th inch) and a wall thickness of 1.3 mm (0.050 inch) connected at one end to a high pressure pump (Hewlett Packard Model HP 1050 TI) and at another end to a back-pressure control device. Between the two ends, the section of tubing was

coiled about a torus-shaped metal mandrel. The mandrel was situated above a primary coil of a transformer so that the coils of tubing and the mandrel functioned as secondary coils of the transformer. The coils of tubing were further equipped with one end of a temperature probe. The other end of the temperature probe was connected to a temperature controlling device. The temperature controlling device regulated the current supplied to the primary coil of the transformer which regulated the heat of inductance imparted to the coiled steel tubing.

[0067] A reaction mixture was prepared by mixing monomers (at a concentration of 50%) in, solvent (acetone) and initiator (2% of tert-butyl peroxide by weight of monomers) at a pressure of 3,500 psi and under the conditions described in Table 1 below. Nitrogen was bubbled through the mixture while stirring. Solvent was pumped through the tubing via the high pressure pump at a rate of from 0.05 to 10 milliliters per minute (ml/min). The pressure was maintained at a level of from 20 MPa (200 bars) to 35 MPa (350 bars). Current was supplied to the primary coil of the transformer to raise the mandrel temperature to 230°. After about 15 minutes, the solvent being pumped through the tubing was replaced by the reaction mixture which was continuously pumped through the tubing at the same rate, temperature and pressure. After allowing the solvent to be cleared from the tubing, product was collected as the effluent from the back-pressure control device. When the addition of the reaction mixture was nearly complete, solvent was pumped through the tubing at the same rate, pressure and temperature as the reaction mixture. Solvent and residual monomer was removed on a rotary evaporator or a wiped-film evaporator.

Table 1 Synthesis conditions

Example	Monomer wt ratio	Reaction temperature / °C
A.1	99 OA / 121 DA / 80 HBA	230°C
A.2	68 BA / 32 HEA	220°C
A.3	63 BA / 37 HBA	210°C
A.4	67 BA / 33 GMA	200°C
A.5	77 LA / 23 GMA	210°C
A.6	32 OA / 45 DA / 23 HEA	235°C
A.7	52 BA / 48 HEA	350°C

Table 2 Product characteristics

Example	Mole ratio	M _n	M _w
A.1*	3.8 OA / 4.6 DA / 4.2 HBA	2200	5600
A.2	1.86 BA / 1.00 HEA	1642	5669
A.3	2.45 BA / 1.00 HBA	1893	6912
A.4	2.07 BA / 1.00 GMA	209	6890
A.5	1.92 LA / 1.00 GMA	2216	5806
A.6	0.93 OA / 1.12 DA / 1.00 HEA	-	-
A.7	1.33 BA / 1.00 HEA	805	2188

* A.1 was a macromonomer which was a liquid at room temperature with a polydispersity of 2.6 as measured by GPC and where approximately 70 % of the macromonomer contained a terminal unsaturated group.

B. Preparation of Nitrogen-functional Macromonomers (Oligomers B.1 & B.2)

[0068] Nitrogen-functional macromonomers of type B are prepared as described below. These monomers may be used as precursors to prepare type C semi-telechelic functional oligomers as described herein and/or may be used directly to prepare formulations of the present invention.

[0069] Oligomer B.1:

A mixture containing 500 g of Oligomer A.1 and 24.9 g of APP, was first heated for 3 hours at 80°C, then heated for 10 hours at 90°C. The relative conversion was followed using proton NMR integration of the C=C-H protons with respect to the –OCH₂- protons of the precursor macromonomer. After heating for another 48 hours at 120°C, the integration ratio indicated that 56 % of Oligomer A.1 was converted to the nitrogen-functional Oligomer B.1. Next, another 6.2 g of APP was added to the mixture. The mixture was further heated for 24 hours at 120°C. The yield of the nitrogen-functional Oligomer B.1 was approximately 60 %. The mixture was passed through a wiped film evaporator to remove un-reacted amine and other impurities to yield a brown liquid.

[0070] Oligomer B.2:

A mixture containing 350 g of Oligomer A.1 and 12.5 g of DMAPA was first heated for 3 hours at 80°C, then heated for 18 hours at 90°C, followed by further heating for 24 hours at 120°C. At this point, the conversion, as measured above for Oligomer B.1, was negligible. To the mixture, 50 g of isopropanol was added and the mixture was then heated for 72 hours at 120°C. After this reaction period, 73 % of Oligomer A.1 was converted to the Oligomer B.2. A second portion of 3.1 g of DMAPA was added to the mixture and the mixture was heated for 24 hours at 120°C. The yield of the nitrogen-functional Oligomer B.2 was approximately 80 %. The mixture was passed through a wiped film evaporator to remove unreacted amine and other impurities to yield a light brown liquid.

C. Preparation of Semi-telechelic nitrogen functional oligomer (Oligomers C.1 & C.2)

[0071] Semi-telechelic functional oligomers may be prepared as described herein and may be used directly to prepare formulations of the present invention.

[0072] A reaction mixture of a Monomer B, AA, HQ, and toluene was added to a 1-liter, 4 neck round bottom flask equipped with a Dean-Stark trap and condenser, a thermometer, an air sparge, and overhead blade stirrer, and a serum stopper. The reaction mixture was heated to reflux at 110°C to 120°C for 30 minutes to removed residual moisture. Next, the catalyst, MSA, was added to the reaction mixture and the reaction mixture was heated at reflux for a period of 5 to 8 hours. The progress of the reaction was monitored by measuring the water removed from the reaction mixture. After no further water was removed, the reaction was stopped by cooling the reaction mixture containing the Oligomer C to room temperature.

Table 3 Reaction Mixtures and Catalyst

Product	Oligomer C.1	Oligomer C.2
Monomer B	348 g of B..1	288 g of B.2
AA	91 g	76 g
Toluene	246 g	288 g
HQ	0.7 g	0.6 g
MSA	5.1 g	14.4 g

[0073] Oligomer C was purified by first mixing the reaction mixture with pre-dried Amberlite IRN-78 ion exchange resin (Amberlite is a trademark of Rohm and Haas Co.) for 1 hour to remove the MSA. The ion exchange resin was removed by filtration and rinsed with toluene and then Oligomer C was passed through a

wiped film evaporator at 100 °C to 120°C and 13.3 kPa (100 mm of Hg) to remove the toluene solvent. Residual AA was removed with a second pass through the wiped film evaporator at 130°C and 0.27 kPa (2 mm of Hg). A titration of Oligomer C showed the nitrogen-functionality had partially complexed with acid. The Oligomer C was dissolved in toluene and treated with Amberlite IRN-78 ion exchange resin to removed the acid complex. After treatment, approximately 500 ppm of HQ was added and the toluene was distilled from Oligomer C. The average composition of each of the Oligomers C.1 and C.2 was calculated using NMR to determine the ratios of the residuals of the first monomer, second monomers, the nitrogen functionality, and GPC to determine the molecular weight. Based on Mn, the weight average molecular weight, the average composition was calculated and is reported in Table 4. The BDDA residue represents the HBA residue which has been esterified with pendant AA. The nitrogen functionality is reported as the starting amine.

Table 4 Average Composition of Oligomers C

	<u>Oligomer C.1</u>	<u>Oligomer C.2</u>
Av. Compn /	3.1 OA / 3.8 DA /	2.6 OA / 3.1 DA /
(wt ratios)	2.2 BDDA / 0.22 APA	2.0 BDDA / 0.37 DMAPA
M _w / daltons	7646	13,702
M _n / daltons	1820	1553
HQ	468 ppm	508 ppm

D. Preparation of Oligomers containing pendant acrylate groups (D.1 to D.4)

[0074] Oligomers with pendant acrylate groups may be prepared as described below (with reference to Table 5A & 5B) and may be used directly to prepare formulations of the present invention.

Reaction

[0075] A reaction flask (i) fitted with an overhead blade stirrer, a thermocouple and temperature controller connecting to a pneumatic jack with heating mantle for heating, a Dean-Stark trap and condenser and an air sparge line was charged with (ii) g of an Oligomer A, (iii) g of AA, (iv) g of toluene and (v) g of HQ. The Dean-Stark trap was filled with toluene and the reaction was heated to reflux (116°C) whilst maintaining a slow stream of air through the batch. After refluxing for about (vi) minutes and removing residual water in the Dean-Stark trap (vii) ml of MSA was added with a syringe. The air sparge and reflux was continued as water was removed and monitored in the Dean-Stark trap. After (viii) hours, (ix) % of the theoretical water was collected and the reaction mixture was cooled. The batch was treated with (x) g of Amberlite IRN-78 for 30 minutes to remove the residual MSA and then filtered. The product was isolated using a wiped film evaporator with the final pass at (xi) °C and about 2 mm / Hg pressure and analysed as described below.

Table 5A – Reaction conditions

Example	(i) – flask	(ii) Olg. A	(iii) - AA	(iv) – Tol.	(v) - HQ
1a = D1	3l, 4-neck	651 g, A2	504 g	690 g	1.86 g
1c = D2	2l, 4-neck	477 g, A3	288 g	394 g	1.16 g
1v =D3	3l, 4-neck	812 g, A6	432 g	768 g	2.0 g

Table 5B – Reaction conditions (continued)

(vi) - reflux time before MSA	(vii) - MSA	(viii) - reflux time with MSA	(ix) - % water	(x) - IRN 78	(xi)- evp. temp.
15 mins	2.2 ml (3.3 g)	6 hrs	103%	30 g	100°C
15 mins	1.3 ml (1.9 g)	5 hrs	124%	20 g	100°C
30 mins	1.9 ml (2.9 g) ¹	3.5 hrs	110%	52 g	130°C

1 At about 30% completion an additional 1.9 ml (2.9 g) of MSA was added.

Products

[0076] Oligomer D1 (Acrylate functional BA/ HEA / AA oligomer prepared from Oligomer A2)

The product prepared and isolated as described above was analysed and found to comprise 0.8% residual AA and 2.1% AA dimer. The batch was dissolved in about an equal volume of toluene and treated with 327 g of Amberlite IRN-78 (washed with methanol and then toluene) for one hour and then filtered. HQ (0.28 g) was added and the sample was concentrated on a Rotovap (60°C with an air sparge) and finally isolated using a wiped film evaporator to give 474 g of the product. The product composition was determined to be (mole ratio by mnr) 2.15 BA / 0.09 HEA / 0.91 HEA -AA ester) and its molecular weight was determined by GPC to be 2,148 (M_n) and 9,300 (M_w).

[0077] Oligomer D2 (Acrylate functional BA/ HBA / AA oligomer prepared from Oligomer A3)

The product prepared and isolated as described above was analysed and found to comprise 0.04 mmol / g of titratable acid. The batch was dissolved in about an equal volume of toluene and treated with 182 g of Amberlite IRN-78 (washed with methanol and then toluene) for one hour and then filtered. HQ (0.23 g) was added and the sample was concentrated on a Rotovap (60°C with an air sparge) and finally isolated using a wiped film evaporator to give 403 g of the product. The product composition was determined to be (mole ratio by mnr) 2.53 BA / 1.00 (HBA -AA ester) and its molecular weight was determined by GPC to be 2,136 (M_n) and 9,698 (M_w).

[0078] Oligomer D3 (Acrylate functional OA / DA / HEA / AA oligomer prepared from Oligomer A6)

The product prepared and isolated as described above and then HQ (0.375 g) was added to 723 g of the product (519 ppm). The product composition was determined to be (mole ratio by mnr) 0.87 OA / 1.05 DA / 0.12 HEA / 0.88 HEA - AA ester) and its molecular weight was determined by GPC to be 2,304 (M_n) and 7,919 (M_w).

[0079] The following formulations with type C Oligomers are known from co-pending application USSN 60/215,721 and are hereby disclaimed.

Oligomer C in known pigment dispersions

[0080] It is known (from USSN 60/215,721) to prepare pigment dispersions using certain Oligomer C type oligomers using pigment slurries and dispersions of oligomer C as follows.

[0081] A pigment slurry by adding 5 g pigment (Black -Raven 450; Irgalite® Blue GLO; Magenta- Irgalite® Rubine L4BD; or Irgalite® Yellow BAW.) and 95 g xylene to a shot mill and agitating the mixture for 20 minutes on a paint shaker. The pigment slurries were filtered through coarse paint filters and stored in sealed bottles. A dispersion was prepared by dissolving 10 g of the Oligomer C in 90 g xylene.

[0082] To a small vial, 100 µl pigment slurry (shaken to re-suspend any settled pigment), 100 µl dispersant solution and 500 µl xylene were added. The vial was capped and sonicated for 10 minutes in an ultrasonic bath to form samples of dispersed pigment which were evaluated as follows. One drop of the dispersed pigment was placed onto a microscope slide, covered with a cover slip, and allowed to sit for 2 minutes. The sample on the microscope slide was observed with a microscope at a magnification of 250x. Flocculation was readily seen as large irregularly shaped agglomerates of particles. Well dispersed samples had a uniform distribution of particles without agglomerates. Samples were rated on a scale of 1 to 5 in which 1 represents a highly agglomerated sample while 5 represents a well dispersed sample.

Table 6 Evaluations of Dispersed Pigments

Dispersant	Cyan	Magenta	Yellow	Black
Oligomer C.1	1	5	2	5
Oligomer C.2	1	5	5	5
Ebecryl 450	3	4	3	1

[0083] Ebecryl 450 (UCB Chemicals) was used as a comparison as a well known pigment dispersant. The results in Table 6 show that it is known that Oligomers C.1 and C.2 are good pigment dispersants for magenta and black pigments and Oligomer C.2 is also a good pigment dispersant for yellow pigments.

[0084] Oligomer C.2 in a conventional radiation curable formulation

It is also known (from USSN 60/215,721) that Oligomer C2 can be added to a conventional radiation curable formulation as follows.

[0085] A pigmented radiation curable formulation was prepared by mixing 10.0 g of Oligomer C.2, 3.9 g Irgalite Rubine L4BD pigment, 10.0 g trimethylolpropane triacrylate (TMPTA) and 1.0 g Irgacure 1173 photoinitiator (Irgacure is a trademark of Ciba Specialty Chemicals). A coated sample was prepared by applying a 25 µm thick film of the formulation onto a glass slide using a doctor blade. The coated sample was exposed to a dose of 1296 mJ/cm² in a Fusion Systems UV processor (Fusion UV Systems Inc.) equipped with a H bulb. The coated sample containing Oligomer C.2 cured to a solid tack-free film, indicating that oligomer was polymerized.

[0086] However such a generic formulation does not have the particularly advantageous properties as claimed for the formulations of the present invention, for example the known formulation is not substantially free of VOC.

[0087] Non limiting examples of the formulations of the inventions will now be described which use one or more oligomers prepared as described herein. The following ingredients are used herein:

[0088] A photoinitiator available commercially from Ciba additives under the trademark "Irgacure® 1173".

[0089] A photoinitiator available commercially from Ciba additives under the trademark "Irgacure® 819".

[0090] 2-Benzyl-2-(dimethylamino)-1-[4-(4-morpholinyl) phenyl]-1-butanone (CAS no. 119313-12-1) which is a photo-initiator available commercially from Ciba additives as a yellow solid under the trademark "Irgacure® 369".

[0091] Ethyl 4-dimethylaminobenzoate (CAS no. 10287-53-3) which is a photo-initiator available commercially from Great Lakes Fine Chemicals as a white solid under the trademark "Quanticure® EPD".

[0092] 2-Isopropylthioxanthone (CAS no. 5495-84-1) which is a photo-initiator available commercially from Great Lakes Fine Chemicals as a yellow solid under the trade name "Quanticure ITX".

[0093] TMPEOTA which denotes trimethylolpropane ethoxy triacrylate (CAS no. 28961-43-5) available commercially from UCB Chemicals.

[0094] DPGDA which denotes dipropylene glycol diacrylate (CAS no. 57472-68-1) available commercially from UCB Chemicals.

[0095] MEHQ which denotes methyl hydroquinone (CAS no. 150-76-5), an inhibitor available commercially from Aldrich Chemicals.

[0096] The oligomer which is available commercially from UCB Chemicals under the trademark Ebecryl® 450.

[0097] The oligomer which is available commercially from UCB Chemicals under the trademark Ebecryl® 140.

[0098] Polyester acrylate oligomer grinding vehicle for flexographic ink which is available commercially from UCB Chemicals under the trade mark Ebecryl® 812.

[0099] Carbon black pigment available commercially from Columbian Chemicals under the tradename Raven 450.

[0100] Copper phthalocyanine colorant available commercially from Ciba Speciality Chemicals, pigments division, under the trademark Irgalite® Blue GLO.

[0101] Magenta colorant available commercially from Ciba Speciality Chemicals, pigments division, under the trademark Irgalite® Rubine L4BD.

[0102] Yellow colorant available commercially from Ciba Speciality Chemicals, pigments division under the trademark Irgalite® Yellow BAW.

Ink Preparation

[0103] The flexographic inks were prepared in two steps. The first step was preparation of the concentrated pigment paste (grind stage) using the Oligomer (70% by weight), and dry pigment (30% by weight). The pigments used were: Magenta - Irgalite® Rubine L4BD, Blue- Irgalite® Blue GLO, Yellow -Irgalite® Yellow BAW; or Black - Raven 450. The Anthony 5x12 three-roll mill was used to grind the pigment and oligomer. After no scratches were observed on the Hegman grind gage, the pigment concentrate was blended 50/50 with a standard letdown vehicle (of acrylated monomers, photoinitiator, and inhibitor – see Table 7) and dispersed for fifteen minutes to produce a final ink.

Table 7 - Ink formulation

<u>Ingredient</u>	<u>%</u>
Concentrate	50.0
TMPEOTA	29.95
DPGDA	12.0
Irgacure® 369	2.0
Quantacure® EPD	2.0
ITX	2.0
MEHQ	0.05
Total	100.0

Table 8
Summary of oligomers used in Inks 1 to 9 of the invention

Ex	Olgmr group	Hydrophobic Hydroxy acrylate	Target Mol Ratio	Degree of Polymn	or Amine	Acrylation group
Ink 1	D1	BA	HEA	2:1	15	NA AA
Ink 2	D2	BA	HBA	2:1	15	NA AA
Ink 3	A4	BA	GMA	2:1	15	NA AA
Ink 4	A5	LA	GMA	2:1	15	NA AA
Ink 5	A1	ODA	HBA	2:1	12	NA AA
Ink 6	C1	ODA	HBA	2:1	12	APP AA
Ink 7	C2	ODA	HBA	2:1	12	DMPA AA
Ink 8	D3	ODA	HEA	2:1	-	NA AA
Ink 9	A7	BA	HEA	1:1	6	NA RP

Footnotes

- is unknown or not measured; NA = Not Applicable ; and "RP" denotes the reaction product of Empol 1022, DiTMP, and AA

[0104] Rheology Tests

All rheology measurements were performed on a Haake RS150 Cone and Plate rheometer equipped with an aluminum 60mm, 1-degree spindle. The rheometer was thermostated by the use of a Peltier device at 25°C for all measurements. The test procedure (Total 20 minutes) was as follows. Hold instrument at 25°C for 300 seconds. Yield point was measured under controlled stress from 0 Pa to 25 Pa at 25°C for 540 seconds. The viscosity curve was measured at controlled rate from 0.06 seconds⁻¹ to 100 seconds⁻¹ at 25°C for 360 seconds.

[0105] Yield Point

A simple definition for the yield point is the smallest possible stress necessary to induce stationary flow of the sample. That means that the sample behaves like a

solid if stresses below the yield point are applied. Structures within the material are not destroyed, the deformation due to the applied stress is reversible, and the shape of the sample regains after setting the applied stress to zero. With stresses larger than the yield point, the structures within the samples are destroyed and the deformation increases dramatically leading to stationary flow of the sample. The transition between the solid and the liquid behaviour can be interpreted as the yield point.

[0106] Viscosity Curve

The measurement of the viscosity curve helps to classify a substance according to one of the possible characteristics: Newtonian, Psuedoplastic, and Thixotropic (see graphs below). With the steady state flow curve in the controlled rate mode, a certain shear rate is selected. Once the steady state conditions have stabilised, a data point for the viscosity will be collected. Then the selected shear rate is increased or decreased in order to obtain a constant shear stress value after a certain time. The shear rate data points are collected from 0 to 100 (1/seconds) to obtain a steady state flow curve. The data reported in the Tables are for shear rates of 50 and 100 (1/seconds).

[0107] Results

The data in Table 9 below shows that incorporating hydrophobic groups into radiation curable acrylated oligomers gives improved pigment wetting for use in for example graphic art applications.

Table 9
Rheology data for Magenta inks of the invention

Ex	Olgmr	Yield	Viscosity	Viscosity	Neat	Resin
		Viscosity	/ Pa	@ 1/50 s.cP	@ 1/100 s.cP	@ 25°C
Ink 1	D1	> 25	3620	2620	29500	
Ink 2	D2	<1	3540	2620	7400	
Ink 3	A4	19	1160	881	760 (@ 60°C)	
Ink 4	A5	>25	1940	1370	1000 (@ 60°C)	
Ink 5	A1	0	1380	990	-	
Ink 8	D3	15	1300	1000	-	

[0108]Comparing Ink 8 (D3) with Ink 1 (D1) and Ink 5 (A1) with Ink 2 (D2) indicates that substitution of the more hydrophobic group ODA for BA gave inks with lower overall viscosities and lower yield points. Both of these attributes are important to achieving inks with the preferred improved performance.

[0109]The data in Table 10 below shows that incorporating of amines or amides into radiation curable acrylated oligomers also gives improved pigment wetting for use in for example graphic art applications.

Table 10
Rheology data for Yellow inks of the invention

Ex	Olgmr(s)	Yield(Pa)	Viscosity (@ 50 s ⁻¹)	Viscosity (@ 100s ⁻¹)
Ink 5	A1	19	3100	2220
Ink 6	C1 & Eb 140	15	3013	2236

[0110] The inks were used to printed samples with a Cavanaugh Hand Proffer. It was observed that the ink of Example 6 which also incorporated Ebecryl 140 (from UCB) exhibited improved transparency and strength when compared to the ink of Example 5. This indicates the amine modification improved the pigment wetting of this yellow pigment.

[0111] The data in Tables 11 to 14 below show that irrespective of pigment colour the inks of the present invention give improved pigment wetting compared to known inks and thus the oligomers used and described herein may act as universal radiation curable pigment wetters. Comp X and Comp Y are comparative inks prepared using the ink formulation of Table 7 using the known oligomers Ebecryl 450 and Ebecryl 812 both available commercially from UCB Chemicals.

Table 11 - Rheology data for Yellow inks

Ex	Oligomer	Yield (Pa)	Viscosity (@ 50 s ⁻¹)	Viscosity (@ 100s ⁻¹)
Comp X	EB 450	5	2,052	1,539
Comp Y	EB 812	8	960	787
Ink 9	A7	5	2,825	2,226

Table 12 - Rheology data for Magenta inks

Ex	Oligomer	Yield (Pa)	Viscosity (@ 50 s ⁻¹)	Viscosity (@ 100s ⁻¹)
Comp X	EB 450	< 1	1,596	1,258
Comp Y	EB 812	20	1,512	1,163
Ink 9	A7	0	2,796	2,681

Table 13 - Rheology data for Cyan inks

Ex	Oligomer	Yield (Pa)	Viscosity (@ 50 s ⁻¹)	Viscosity (@ 100s ⁻¹)
Comp X	EB 450	17	1,256	920
Comp Y	EB 812	11	1,008	821
Ink 9	A7	10	1,480	1,210

Table 14 - Rheology data for Black inks

Ex	Oligomer	Yield (Pa)	Viscosity (@ 50 s ⁻¹)	Viscosity (@ 100s ⁻¹)
Comp X	EB 450	>25	1,616	1,023
Comp Y	EB 812	21	1,325	937
Ink 9	A7	15	1,540	1,290

[0112] Thus an ink of the invention (Ink 9) compares favorably with known inks prepared from two commercially available resins. In the case of Comp X, yield points for black and cyan are very high and will result in an ink that prints poorly on press. Comp Y has large yield points for black and magenta inks compared to Ink 9 which demonstrates consistently low yield points for all colors.